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### Erratum

## Erratum to “Differentiation of pancreatic endocrine progenitors reversibly blocked by premature induction of MafA” [Dev. Biol. 385 (2014) 2–12]



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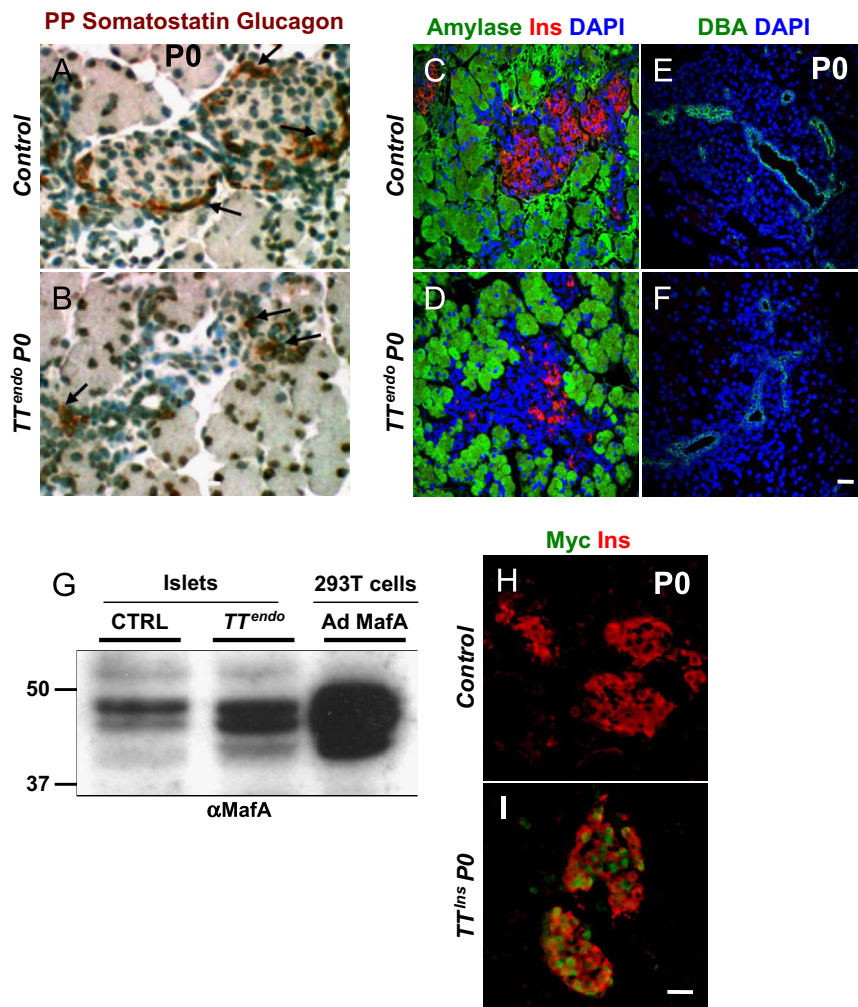
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Fig. 2I shows an image from  $TT^{ins}PO$  pancreas and not  $TT^{endo}PO$  as indicated in the published article. The correct figure and legend are provided here.

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**Fig. 2.** Mistimed MafA expression in endocrine progenitors does not alter fate selection. (A and B) P0 pancreatic sections immunostained with antibody cocktail against glucagon, somatostatin and PP (brown, peroxidase staining) showing an overall reduction in hormone<sup>+</sup> cells in *TT<sup>endo</sup>P0* compared to control littermates. Magnification bar = 20 μm. (C–F) Amylase<sup>+</sup> and DBA<sup>+</sup> duct (green) areas were not affected in *TT<sup>endo</sup>P0* pancreas that has reduced insulin<sup>+</sup> (red) cells in islet-like structure. (G–I) Transgene expression level was not detrimental to endocrine cells. (G) Western blot analysis of extracts from *TT<sup>endo</sup>* islets isolated from adult mice sacrificed after receiving DOX for 5 days show a 2–4 fold increase in total MafA protein compared to extracts from comparable control islets; 293T cells infected with Adeno-MafA (AdMafA) virus as positive control. (H–I) Induction of MafA<sup>Myc</sup> (green) expression in insulin<sup>+</sup> (red) cells in *TT<sup>Ins</sup>* (*Ins<sup>CRE</sup>*; *Rosa26<sup>rtTA</sup>*; *TetO<sup>MafA</sup>*) animals receiving DOX from E7.5 until sacrifice at P0 did not affect the formation of insulin<sup>+</sup> cells. Images of control animals in A, C, and E were from double transgenic *Ngn3<sup>Cre/+</sup>*; *Rosa26<sup>rtTA/+</sup>*, while those for H were from double transgenic *Ins<sup>Cre/+</sup>*; *Rosa26<sup>rtTA/+</sup>*. Magnification bar = 20 μm.